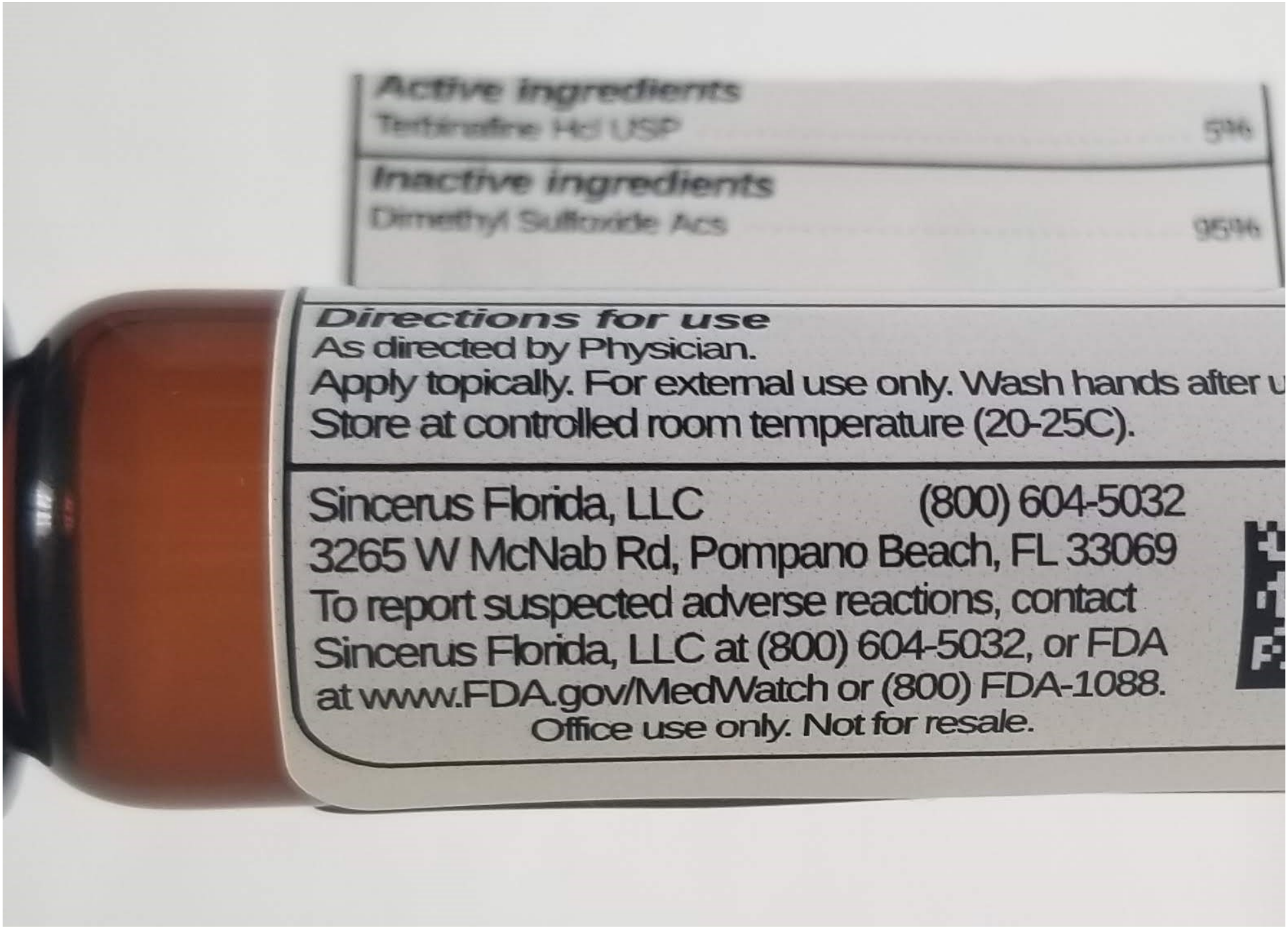
TERBINAFINE HCL 5%- terbinafine hcl 5% solution
Sincerus Florida, LLC

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

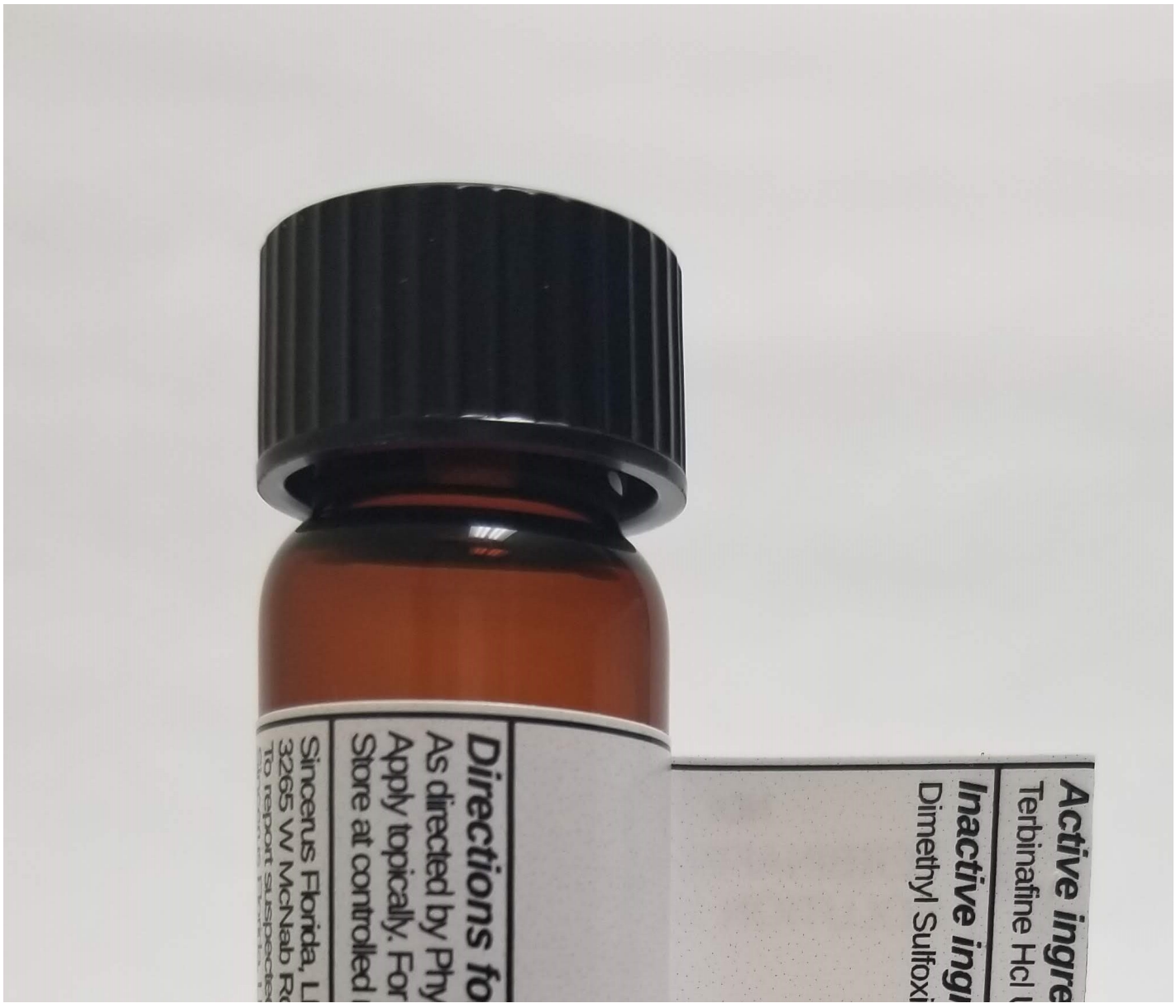
TERBINAFINE HCL 5%

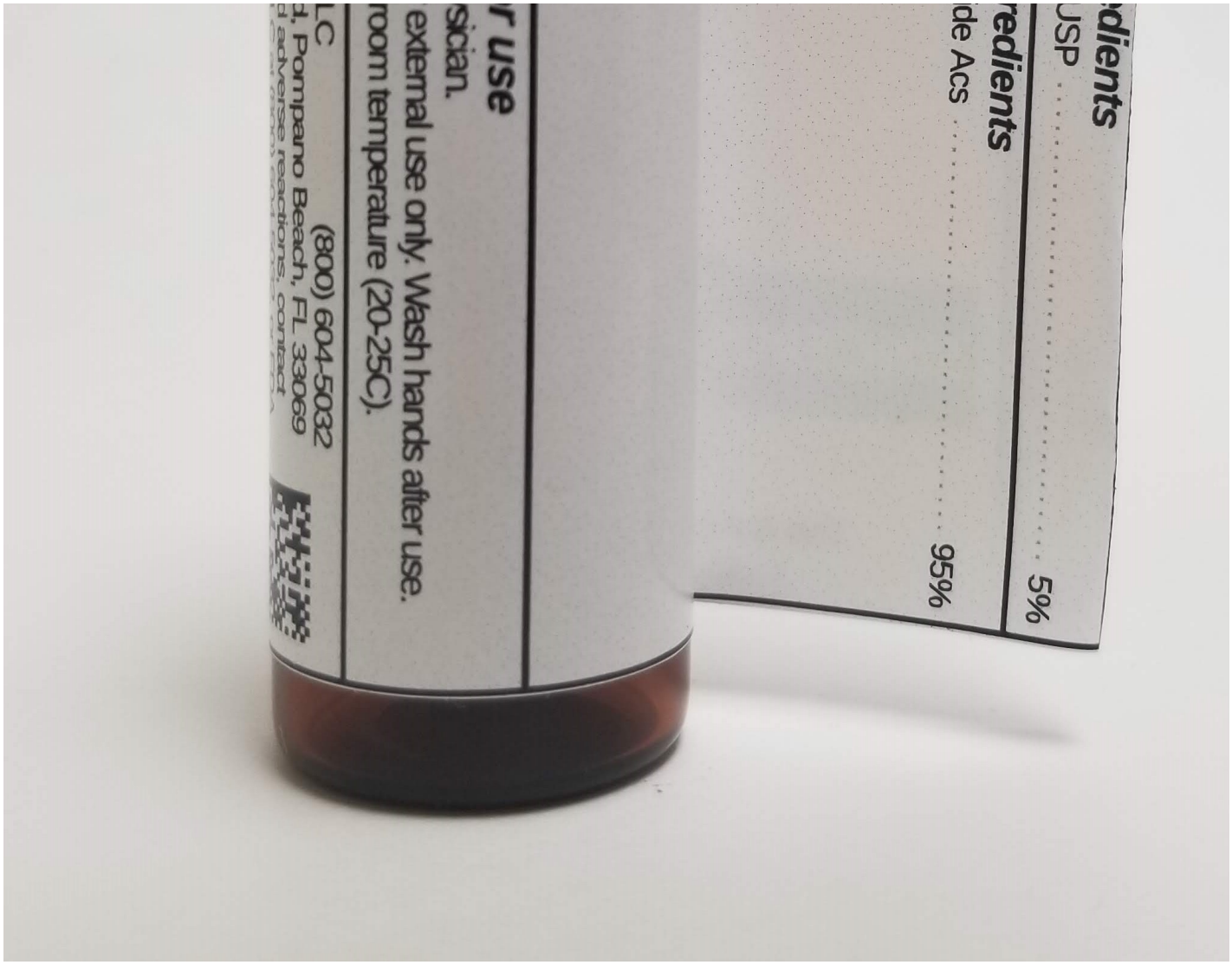
Directions for use

Sincerus Florida, LLC. Adverse reactions

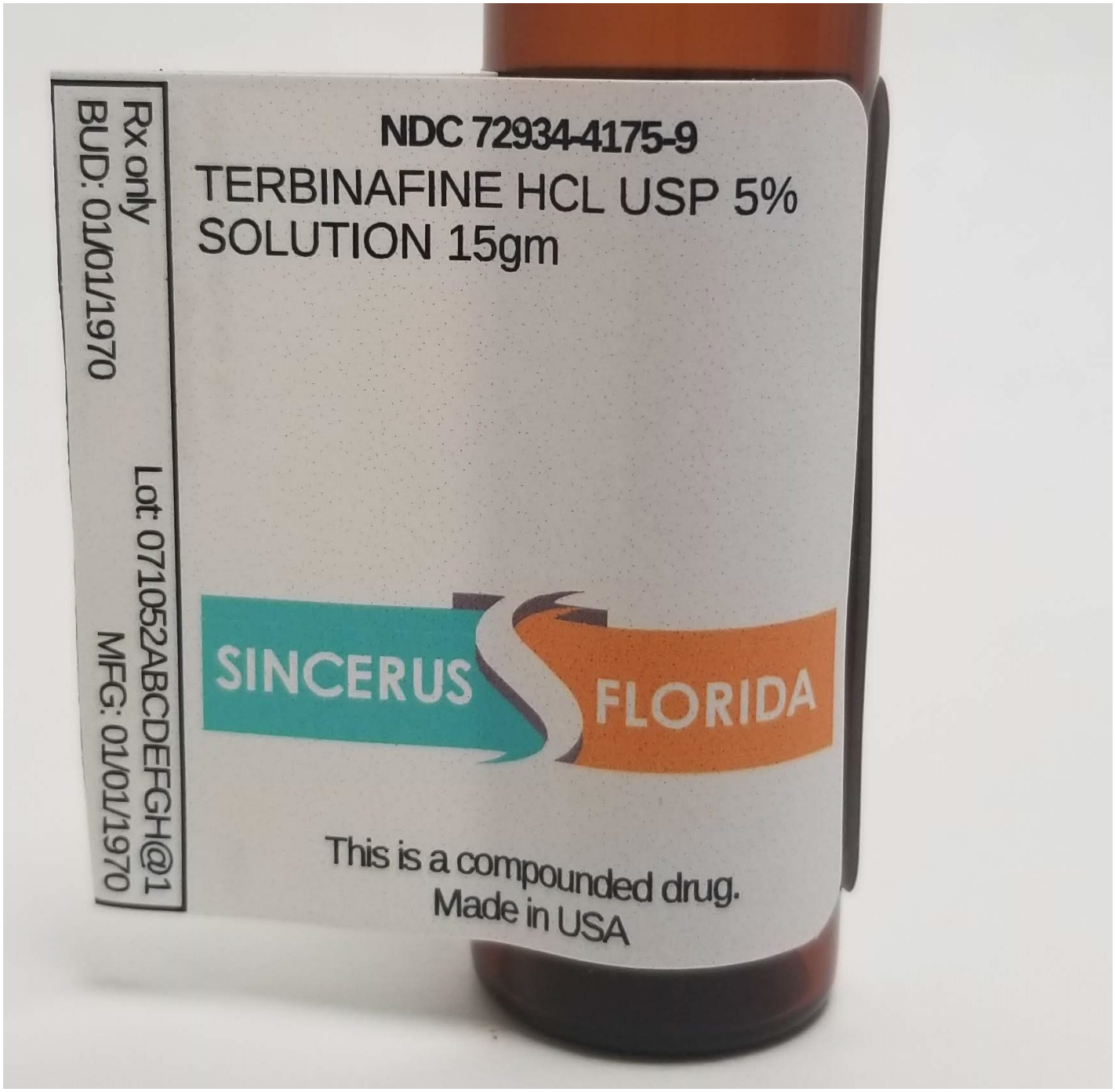

Active ingredients
Terbinafine Hcl USP 5%
Inactive ingredients
Dimethyl Sulfoxide Acs 95%
Directions for use
As directed by Physician.
Apply topically. For external use only. Wash hands after u
Store at controlled room temperature (20-25C).
Sincerus Florida, LLC (800) 604-5032
3265 W McNab Rd, Pompano Beach, FL 33069
To report suspected adverse reactions, contact
Sincerus Florida, LLC at (800) 604-5032, or FDA
at www.FDA.gov/MedWatch or (800) FDA-1088.
Office use only. Not for resale.

Active, inactive





NDC 72934- 4175-9 TERBINAFINE HCL USP 5%. Solution 15 gm



TERBINAFINE HCL 5%

terbinafine hcl 5% solution

Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	**Item Code (Source)**	NDC:72934-4175
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
TERBINAFINE HYDROCHLORIDE (UNII: 012C11ZU6G) (TERBINAFINE - UNII:G7RIW8S0XP)	TERBINAFINE HYDROCHLORIDE	5 g in 100 g

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72934-4175-9	15 g in 1 VIAL; Type 0: Not a Combination Product	05/21/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		05/21/2019	

Labeler - Sincerus Florida, LLC (080105003)

Establishment

Name	Address	ID/FEI	Business Operations
Sincerus Florida, LLC		080105003	manufacture(72934-4175)

Revised: 5/2019 Sincerus Florida, LLC